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Exhibit 2

510K Summary: K072532

DOOYANG SYSTEMS INC.

SK Techno Bldg. 7th Floor 16-4, Sungsoo 1-Ga, 2-Dong, Sungdong-Ku, Seoul, 133-710 Korea

TEL : 82-2-489-2612

FAX : 82-2-467-4915

Contact: Yun-Hwoi, Ku /Managing Director

Date : October 30, 2007

1. Identification of Device :

   Proprietary-Trade Name: "Truepix LCD Medical Display Monitor" (Models : TX-5M, TX-3M, TX-2M)

   Classification Name: System, Image Processing, Radiological, Product Code: LLZ

   Common/Usual Name: Radiological Image Processing System, LCD Monitor

2. Equivalent Legally Marketed Devices:

   Lumimed MM20, MM30 and MM50 Monitors, K052120

3. Indication for Use (Intended Use) :

   Monochrome LCD Monitors (TX-5M, TX-3M, TX-2M) intended to be used in various kinds of medical image application for which the device complies with the performance specified by the manufacturer of the system. Not for use in mammography.

4. Description of the Device

   Monochrome LCD Monitors(TX-5M, TX-3M, TX-2M) are used to display images such as X-ray, MRI images. These models have resolutions of: 2560x2048, 2048x1536, 1600x1200. These models have USB functions and an optional photo sensor GFU12-SEQ01 made by SEQUEL IMAGING INC. These models are certified to the EN60601-1 medical safety standard. The monitors use universal Power Supply compatible with AC100V~240V, 50/60Hz. The graphics card needed for the personal computer is the Real Vision & Matrox Series.
   The monitors may be deployed in the portrait or the landscape position.

5. Safety and Effectiveness, comparison to predicate device

Items	Lumimed MM20, MM30 and MM50 Monitors,	New		
		Truepix TX-2M	Truepix TX-3M	Truepix TX-5M
510(k) Numbers	K052120	New		
Manufactures	Heeyoung CI.	Dooynag Systems Inc.		
Panel Size and Type	21.3" /20.1" TFT monochrome LCD display	21.3" TFT monochrome LCD display	21.3" TFT monochrome LCD display	20.1" TFT monochrome LCD display
Pixel Pitch	0.270mm x 0.270mm 0.2115mm x 0.2115mm 0.156mm x 0.156mm	0.270mm x 0.270mm	0.2115mm x 0.2115mm	0.156mm x 0.156mm
Available Cabinet Colors	Black	Black	Black	Black
Native Resolutions	1200x1600(portrait) 1600x1200(landscape) 1536x2048(portrait) 2048x1536(landscape) or 2048x2560(portrait) 2560x2048(landscape)	1200x1600 (portrait) 1600x1200 (landscape)	1536x2048 (portrait) 2048x1536 (landscape)	2048x2560 (portrait) 2560x2048 (landscape)
Brightness	1000cd/m2 900cd/m2 850cd/m2	1000cd/m2	800cd/m2	850cd/m2
Contrast Ratio	700:1 700:1 600:1	700:1	700:1	600:1
Dot Clock	125MHz 130MHz 162MHz	129MHz	130MHz	166.52MHz
Network Interface	USB (1 Up, 2 Downstream)	USB (1 Up, 4 Downstream)	USB (1 Up, 4 Downstream)	USB (1 Up, 4 Downstream)
Active Display Size (HxV)	432(H) x 324(V) 432(H) x 324(V) 399(H) x 319(V)	433.152x324.864mm	433.152x324.864mm	399.36x319.49mm
Dimension (WxHxD)	472 x 495 x92 472 x 495 x92 438 x 458 x 98	505 w x 505 h x 81 d	505 w x 505 h x 81 d	445 w x 376 h x 90 d
Luminance Calibration	Software(Optional) Photo-sensor(Optional)	Software(Optional) Photo-sensor(Optional)	Software(Optional) Photo-sensor(Optional)	Software(Optional) Photo-sensor(Optional)
Power	AC 100~240Volts/60Hz DC 12V,6,67A	AC 100~240Volts/60Hz DC 12V,6,67A	AC 100~240Volts/60Hz DC 12V,6,67A	AC 100~240Volts/60Hz DC 12V,6,67A

6. Testing Information and Conclusion

In all material respects, the Truepix LCD Medical Display Monitors are substantially equivalent to the predicate device. Testing was performed according to internal company procedures and the monitors were safety certified to International Standard.

DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

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Dooyang Systems, Inc.
% Mr. Daniel Kamm, P.E.
Regulatory Engineer
Kamm & Associates
PO Box 7007
DEERFIELD IL 60015

Re: K072532
Trade/Device Name: Truepix LCD Medical Display Monitor
Regulation Number: 21 CFR 892.2050
Regulation Name: Picture archiving and communications system
Regulatory Class: II
Product Code: LLZ
Dated: September 4, 2007
Received: September 17, 2007

Dear Mr. Kamm:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

This letter will allow you to begin marketing your device as described in your Section 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801), please contact the Center for Devices and Radiological Health's (CDRH's) Office of Compliance at one of the following numbers, based on the regulation number at the top of this letter.

21 CFR 876.xxxx	(Gastroenterology/Renal/Urology)	240-276-0115
21 CFR 884.xxxx	(Obstetrics/Gynecology)	240-276-0115
21 CFR 892.xxxx	(Radiology)	240-276-0120
Other		240-276-0100

Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR Part 807.97). For questions regarding postmarket surveillance, please contact CDRH's Office of Surveillance and Biometric's (OSB's) Division of Postmarket Surveillance at 240-276-3474. For questions regarding the reporting of device adverse events (Medical Device Reporting (MDR)), please contact the Division of Surveillance Systems at 240-276-3464. You may obtain other general information on your responsibilities under the Act from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (240) 276-3150 or at its Internet address http://www.fda.gov/cdrh/industry/support/index.html.

Sincerely yours,

Nancy C Brogdon

Nancy C. Brogdon
Director, Division of Reproductive,
Abdominal, and Radiological Devices
Office of Device Evaluation
Center for Devices and Radiological Health

Enclosure

Indications for Use

510(k) Number (if known): K072532

Device Name: "Truepix LCD Medical Display Monitor"

Indications For Use: Monochrome LCD Monitors (TX-5M, TX-3M, TX-2M) intended to be used in various kinds of medical image application for which the device complies with the performance specified by the manufacturer of the system. These monitors are not intended for use in mammography.

Prescription Use X AND/OR Over-The-Counter Use _____
(Part 21 CFR 801 Subpart D) (21 CFR 807 Subpart C)

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE IF NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)

(Division Sign-Off)
Division of Reproductive, Abdominal and Radiological Devices
510(k) Number K072532